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(2) determining the presence or absence of an endocrine disrupting action of the test substance by obtaining a first gene expression pattern obtained from

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the cell in the first culture system and a third gene expression pattern obtained from the cell in the third culture system, comparing the first gene expression pattern with the third gene expression pattern and a second expression pattern expressed by a cell having a sensitivity to the endocrine hormone in the presence of the endocrine hormone, thereby detecting a gene specific to the first gene expression pattern.

3. A method of detecting an endocrine disrupting action of a test substance, comprising:

(1) (a) culturing a cell having a sensitivity to an endocrine hormone in a first culture system in which the endocrine hormone and the test substance are present;

(b) culturing the cell a having a sensitivity to the endocrine hormone in a third culture system in which the endocrine hormone is absent and the test substance is present;

(2) determining the presence or absence of an endocrine disrupting action of the test substance by comparing a first gene expression pattern obtained from the cell of the first culture system with a third gene expression pattern obtained from the cell in the third culture system, thereby detecting a gene specific to the first gene expression pattern.

4. A method of detecting an endocrine disrupting action of a test substance, comprising:

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(1) (a) culturing a cell having a sensitivity to an endocrine hormone in a first culture system in which the endocrine hormone and the test substance are present;

5 (b) culturing the cell a having a sensitivity to the endocrine hormone in a second culture system in which the endocrine hormone is present and the test substance is absent; and

10 (c) culturing the cell a having a sensitivity to the endocrine hormone in a third culture system in which the endocrine hormone is absent and the test substance is present; and

15 (2) determining the presence or absence of an endocrine disrupting action of the test substance by comparing a first gene expression pattern obtained from the cell in the first culture system with a second gene expression pattern obtained from the cell in the second culture system and the third gene expression pattern obtained from the cell in the third culture system, thereby detecting a gene specific to the first gene expression pattern.

20 5. The method according to any one of claims 1 to 4, wherein said endocrine hormone is selected from the group consisting of estrogen, estradiol, progesterone, androgen, testosterone, androsterone, 25 cortisol, aldosterone, corticosterone, cortison, triiodothyronine, and tyroxacin; and said cell having

6. The method according to any one of claims 1 to 4, wherein said endocrine hormone is triiodothyronine and said cell having a sensitivity to the endocrine hormone is Neuro2a.

7. The method according to any one of claims 1 to 4, wherein comparison of the gene expression patterns are made by comparing bands obtained by subjecting a gene group contained in each of gene expression patterns to electrophoretic separation.

8. The method of detecting an endocrine disrupting action of a test substance according to any one of claims 1 to 4, comprising

- (a) recovering RNAs from each of the culture systems of (1);
- (b) subjecting the RNAs recovered in the step (a) to reverse transcription;
- (c) amplifying reverse transcription products obtained in (b) by PCR; and
- (d) subjecting PCR products obtained in the step (c) to electrophoresis, comparing electrophoretic patterns of bands obtained, thereby detecting a band specific to a first gene expression pattern.

9. The method according to any one of claims 1 to 4, wherein said gene expression patterns are

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compared by hybridizing gene groups contained in each of the gene expression patterns with each other, and subtracting unhybridized genes.

10. A method of detecting an endocrine disrupting action of a test substance, comprising:

(1) determining the presence or absence of an endocrine disrupting action of the test substance by

(a) culturing a cell having a sensitivity to an endocrine hormone in a first culture system in which the endocrine hormone and the test substance are present;

(b) culturing a cell having a sensitivity to the endocrine hormone in a second culture system in which an endocrine hormone is present and test substance is absent;

(c) culturing a cell having a sensitivity to the endocrine hormone in a third culture system in which the endocrine hormone is absent and the test substance is present; and

(d) culturing a cell having a sensitivity to the endocrine hormone in a fourth culture system in which both the endocrine hormone and the test substance are absent;

(2) (a) isolating RNA from the first culture system and preparing a first cDNA based on the RNA;

(b) isolating a second RNA from the second culture system;

(c) isolating a third RNA from the third culture system; and

(d) isolating RNA from the fourth culture system and preparing a fourth cDNA based on the RNA;

5 (3) (a) hybridizing the first cDNA and the second RNA and recovering unhybridized cDNA;

(b) hybridizing the third RNA and the fourth cDNA, recovering unhybridized RNA; and

10 (4) hybridizing the cDNA obtained in (a) of (3) and the RNA obtained in (b) of (3); and recovering unhybridized RNA, thereby detecting a gene specific to an endocrine disrupting action.

11. A polynucleotide obtained in a method comprising the following, and a complementary
15 polynucleotide having a complementary sequence of said polynucleotide,

(1) (a) culturing a cell having a sensitivity to an endocrine hormone in a first culture system in which the endocrine hormone and a test substance are present;

20 (b) culturing a cell having a sensitivity to the endocrine hormone in a second culture system in which the endocrine hormone is present and the test substance is absent; and

(c) culturing a cell having a sensitivity to
25 the endocrine hormone in a third culture system in which the endocrine hormone is absent and the test substance is present; and

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(2) determining the presence or absence of an endocrine disrupting action of the test substance by

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5 comparing a first glycoprotein pattern obtained from the cell of the first culture system with a second glycoprotein pattern expressed by a cell having a sensitivity to the endocrine hormone in the presence of the endocrine hormone, thereby detecting a glycoprotein specific to the first glycoprotein pattern.

15. A method of detecting an endocrine disrupting action of a test substance, comprising:

10 (1) (a) culturing a cell having a sensitivity to an endocrine hormone in a first culture system with the endocrine hormone and the test substance; and

15 (b) culturing a cell having a sensitivity to the endocrine hormone in a third culture system in which the endocrine hormone is absent and the test substance is present; and

(2) determining the presence or absence of endocrine disrupting action of the test substance by

20 obtaining a first glycoprotein pattern obtained from the cell of the first culture system and a third glycoprotein pattern obtained from the cell of the third culture system, and

25 comparing the first glycoprotein pattern with the third glycoprotein pattern and a second glycoprotein pattern obtained from a cell having a sensitivity to the endocrine hormone in the presence of the endocrine hormone, thereby detecting a glycoprotein specific to the first glycoprotein pattern.

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(b) culturing a cell having a sensitivity to the endocrine hormone in a second culture system in which the endocrine hormone is present and the test

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substance is absent; and

(c) culturing a cell having a sensitivity to the endocrine hormone in a third culture system in which the endocrine hormone is absent and the test substance is present; and

(2) determining the presence or absence of an endocrine disrupting action of the test substance by comparing first, second and third glycoprotein patterns obtained respectively from the cells of the first, second, third culture systems, thereby detecting a glycoprotein specific to the first glycoprotein pattern.

18. The method according to any one of claims 14 to 17, wherein said endocrine hormone is selected from the group consisting of estrogen, estradiol, progesterone, androgen, testosterone, androsterone, cortisol, aldosterone, corticosterone, cortison, triiodothyronine, and tyroxacin; and said cell having a sensitivity to the endocrine disrupting action is selected from the group consisting of Neuro2a, S-20Y, MCF7, TM3, TM4 and 15P-1.

19. The method according to any one of claims 14 to 17, wherein said endocrine hormone is triiodethyronine and said cell having a sensitivity to the endocrine hormone is Neuro2a.

20. The method according to any one of claims 14 to 17, wherein the glycoprotein specific to the first

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21. An antibody against a substance susceptible to an endocrine disrupting action, said antibody being prepared by a method comprising:

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(c) culturing a cell having a sensitivity to the endocrine hormone in a third culture system in which the endocrine hormone is absent and the test

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(3) recovering a glycoprotein by binding a substance capable of specifically binding to a polysaccharide chain contained in the proteins extracted in (2) to the polysaccharide chain;

(5) comparing the glycoprotein patterns to each other, thereby detecting a glycoprotein specific to the first culture system, and further recovering a protein contained in the glycoprotein.

(7) recovering a desired glycoprotein from the proteins extracted in (2) by the polyclonal antibody obtained; and

(8) preparing an antibody against to a polysaccharide chain contained in the glycoprotein obtained in (7).

22. The antibody against the polysaccharide chain susceptible to an endocrine disrupting action,

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